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Amendments To The Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of claims:

(Currently Amended) A pharmaceutical composition comprising:

 a microparticle having an mean particle size of between about 0.51.0μm and about 100
 μm, said microparticle comprising:

a polymeric support material in which a substance can be dispersed, wherein the support material comprises at least about 50% w/w of at least one homopolymer with a repeat unit according to Formula (I):

wherein

R₁ represents a C₁-C₆ alkyl group or a group (CH₂)_m-COOR₃ wherein m is an integer from 1 to 5 and R₃ is a C₁-C₆ alkyl group, R₁ and R₃ being the same or different;

 R_2 represents a $C_1\text{-}C_6$ alkyl group the same or different from R_1 and R_3 ;

n is an integer from 1 to 5; and

at least one therapeutic agent that is encapsulated or dispersed in the polymeric support material of the microparticle.

- 2. (Original) A pharmaceutical composition according to claim 1 wherein: R_1 and R_2 are independently chosen C_1 - C_6 alkyl groups; and n is 1.
- 3. (Original) A pharmaceutical composition according to claim 1 wherein:

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the stated homopolymer comprising repeat units according to Formula (I) wherein R_1 and R_2 are ethyl groups; and

n=1.

- 4. (Original) A pharmaceutical composition according to claim 3, wherein the composition being obtained by a single emulsification process.
- 5. (Currently Amended) A pharmaceutical composition according to any one of claims 1 to 4 wherein the support material comprises:

from about 90 to about 99.5% by weight of a homopolymer as defined in claim 1; and from about 0.5 to about 10% by weight of a polymer additive.

- 6. (Original) A pharmaccutical composition according to claim 5 wherein the polymer additive comprises at least one of polyethyleneoxide, polyvinylalcohol, polyvinylpyrrolidone, poly(N-2-hydroxypropyl methacrylamide), polyhydroxyethylmethacrylate, hydrophilic poly(aminoacid) such as polylysine or polysaccharide.
- 7. (Currently Amended) A pharmaceutical composition according to claim 5 wherein the polymer additive is a polyvinylalcohol.
- 8. (Currently Amended) A pharmaceutical composition according to claim 1 wherein the dispersed substance is hydrophobic.
- (Currently Amended) A pharmaceutical composition according to claim 1 wherein the dispersed substance is a therapeutic agent that requires a solvation vehicle for administration.
- 10. (Currently Amended) A pharmaceutical composition according to claim 1 wherein the dispersed substance is hydrophylic.

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- 11. (Currently Amended) A pharmaceutical composition according to claim 1, wherein the dispersed substance is a therapeutic agent.
- 12. (Withdrawn) A pharmaccutical composition according to claim 1, wherein the dispersed substance is a peptide or polypeptide.
- 13. (Withdrawn) A pharmaceutical composition according to claim 1 wherein the dispersed substance is a protein.
- 14. (Currently Amended) A pharmaceutical composition according to any one of claim 1 wherein the dispersed substance is a bioactive molecule such as a drug, a therapeutic agent, an anticancer agent, a gene therapy agent, a plasmid DNA, a protein, an enzyme, a peptide, a radionuclide, a protein inhibitor, an analgesic, an anti-inflamatory agent, an antibiotic, an antiviral agent, an antineoplastic agent, a pyrimidine, purine or folic acid analog, an cytotoxic agent, an immunomodulator, a hormone, an antibody or a painkiller.
- 15. (Currently Amended) The pharmaceutical composition of Claim 14 wherein the pyrimidine analog is fluorouracil (5-FU).
- 16. (Currently Amended) A pharmaccutical composition according to claim 1 wherein the dispersed substance is a bioactive molecule such as an anticancer agent or a gene therapy agent.
- 17. (Currently Amended) A pharmaccutical composition according to claims 1 wherein the dispersed substance is a therapeutic agent for treating or reducing the severity of a urological disease or disorder.

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- 18. (Currently Amended) A pharmaceutical composition according to claim 1 wherein the dispersed substance is a therapeutic agent for bladder cancer.
- 19. (Currently Amended) A pharmaceutical composition according to any one of claim 1, wherein the dispersed substance is a taxane.
- 20. (Original) A pharmaceutical composition according to claim 19, wherein the taxane is paclitaxel, docetaxel (Taxoterc®) or taxol®.
 - 21-36. (Cancelled).
- 37. (Currently Amended) A method for treating a urological disease or disorder comprising:

administering intravesically a microparticle having a mean particle size of between about 1.0 µm and 100 µm with one or more encapsulated therapeutic agents to the lumen of the bladder-;

contacting the particles to the surface of the mucosa,

releasing the encapsulated therapeutic agent in a controlled manner to treat the urological disease or disorder.

- 38. (Original) A method according to claim 37 wherein the microparticle comprises a poly(methylidene malonate 2.1.2) polymer support material.
- 39. (Original) A method according to claim 37 wherein the urological disorder is a cancer and the microparticle encapsulated therapeutic agent is an anticancer agent.
- 40. (Currently Amended) A method according to claim 37 wherein the anticancer agent is a taxane.

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- 41. (Original) A method according to any to claim 40 wherein the taxane is paclitaxel, docetaxel (Taxotere®) or taxol®.
- 42. (Currently Amended) A method according to claim 37, wherein microparticles with encapsulated paclitaxel are used for intravesical chemotherapy of bladder cancer.

43-51. (Cancelled).

- 52. (Previously Presented) The pharmaceutical composition of claim 1, wherein the microparticle has a mean particle size of between about 1.0 μ m and 20 μ m.
- 53. (New) The method of claim 37, wherein the microparticle has a mean particle size of between about 1.0 μm and 20 μm .